



INVESTOR IN PEOPLE
REC'D 05 OCT 2000

The Patent Office PCT
WIPO

Concept House

Cardiff Road

Newport

South Wales

NP10 8QQ

**PRIORITY
DOCUMENT**

SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH RULE 17.1(a) OR (b)

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

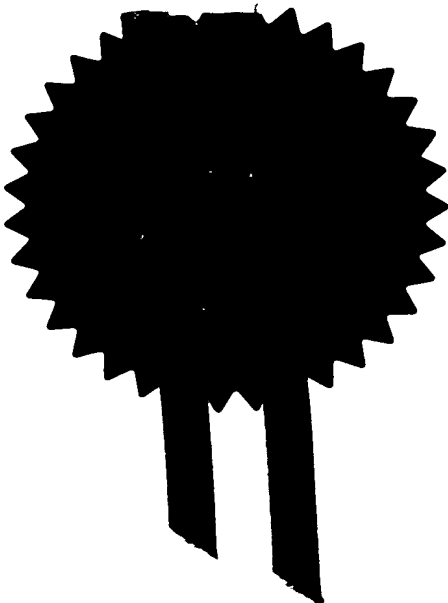
In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

Signed

Andrew Gersey
21 September 2000

Dated 21 September 2000





Patents Form 1/77

Patent Office
Cardiff

THE PATENT OFFICE
20 SEP 1999
RECEIVED BY FAX

1/77
20SEP99 E477902-1 D02944
P01/7700 0.00 - 9922136.8

The Patent Office

Cardiff Road
Newport
Gwent NP9 1RH

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

20 SEP 1999

1. Your reference

SMC 60384

2. Patent application number
(The Patent Office will fill in this part)

9922136.8

3. Full name, address and postcode of the or of each applicant (underline all surnames)

Avecia Limited
Hexagon House
Blackley
Manchester M9 8ZS

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

United Kingdom

4. Title of the invention

Compounds, Compositions and Use

5. Name of your agent (if you have one)

KIRK, Martin John

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

Avecia Limited
Hexagon House
PO Box 42
Blackley
Manchester M9 8ZS

Patents ADP number (if you know it)

7147440001

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)

Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

- a) any applicant named in part 3 is not an inventor, or
 - b) there is an inventor who is not named as an applicant, or
 - c) any named applicant is a corporate body.
- See note (d))

Patents Form 1/77

Patents Form 1/77

THE PATENT OFFICE

20 SEP 1999

RECEIVED BY FAX

9. Enter the number of sheets for each of the following items you are filing with this form. Do not count copies of the same document

Continuation sheets of this form

Description

Claim(s)

Abstract

Drawing(s)

30 ✓

05 ✓

01 ✓

DML

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77)

Request for substantive examination (Patents Form 10/77)

Any other documents (Please specify)

11.

I/We request the grant of a patent on the basis of this application.

Signature K.M. Pinder Date 20/09/99
Avecia Limited Authorised Signatory

12. Name and daytime telephone number of person to contact in the United Kingdom

K.M.Pinder/G.Terry 0161 721 1361/2

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 0645 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- For details of the fee and ways to pay please contact the Patent Office.

Patents Form 1/77

COMPOUNDS, COMPOSITIONS AND USE

This invention relates to colorants, inks and patterned substrates (such as colour filters) comprising these colorants and methods of making the patterned substrates.

The ability to produce brightly coloured patterns or images on substrates, where the colour has high resistance (fastness) to light, water, heat and/or solvents is important in many areas of for instance the electronics and printing industries. Examples of this are in the production of colour filters, in ink-jet printing and in electrophotographic imaging. Thus it is important to discover colorants which can be readily used (and/or formulated into inks for use) in such applications.

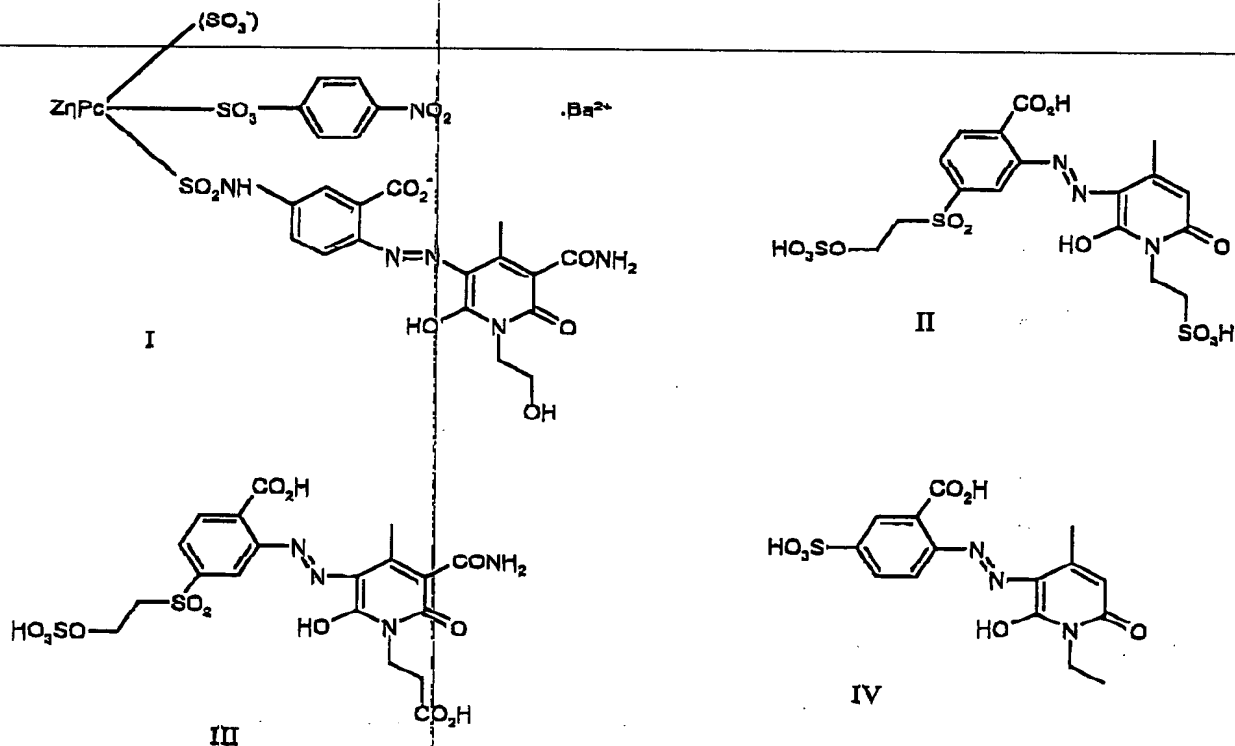
One common form of patterned coloured substrate is a colour filter. Colour filters, alternatively known as optical filters, are a component of coloured liquid crystal displays (LCDs) used as flat screen displays as, for example, in small television receivers or portable computers. Typically, a white back-light is shone through a liquid crystal layer and then a colour filter to produce an image of the desired colour by the transmitted light. The LCD layer comprises an addressable array of pixels. The light at any pixel can be switched on and off by applying a voltage to the liquid crystal film which changes the orientation of the polarising liquid crystals to block the back-light. The pixels are in register with a trichromatic array of colour filter elements to produce a full colour screen capable of displaying images. Some LCD displays are constructed to be viewed by reflected light, but still require a colour filter to produce a full colour image. Colour filters are equally useful for other display technologies such as plasma display panels, cathode ray tubes and electroluminescent displays and as a component of solid state imaging devices. New colorants with properties which are useful in colour filters are desirable.

Many methods can be used to produce colour filters for example: printing, photolithography, laser ablation, electrodeposition, photography and thermal transfer. Of these, printing methods (particularly ink-jet printing [IJP]) and photolithography are preferred. Ink jet printing methods involve a non-impact printing technique for printing an image onto a substrate using ink droplets ejected through a fine nozzle onto a substrate without bringing the nozzle into contact with the substrate. Photolithographic methods involve application to a substrate of a photosensitive ink which can be patterned by selective exposure to radiation (e.g. UV light) through a patterned photomask and subsequent removal of either the exposed or unexposed areas, depending on the type of ink employed.

There are many demanding performance requirements for colorants and inks used in IJP. For example, they desirably provide bright, sharp images having good water-fastness, light-fastness and optical density. The inks are often required to dry quickly when applied to a substrate, but they should not form a crust over the tip of an ink jet nozzle because this will stop the printer from working. The inks should also be stable to

storage over time without decomposing or forming a precipitate which could block the fine nozzle. The most popular ink jet printers are the thermal and piezoelectric ink jet printers. There is a need for colorants and inks which optionally are suitable for both thermal and piezo ink jet printers, have high brightness and colour strength and produce images having a high light-fastness and water-fastness when printed on a substrate.

Dyes of the following formulae are known:



Dyes of structures II & III are water soluble monoazo pyridones used as cellulose reactive dyes as described in EP 169,457 B1 (Hoechst). The dye of structure IV is described in GB 1,359,171 (Ciba-Geigy), as a dye for wool or cellulosics. The dye of structure I in which a pyridone moiety is bound to a zinc phthalocyanine is described in JP-A 01-303407 (Nippon Kayaku) as a one component green dye for colour filters. This is different from the present invention in which a yellow dye is used as a separate component of a green ink for colour filters. This offers much greater flexibility in adjusting the precise shade of the ink and also is advantageous over zinc phthalocyanines which are less favoured due to poor light fastness.

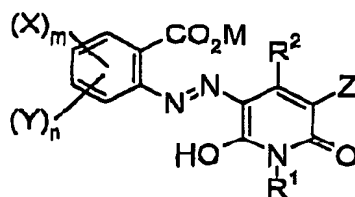
Known mixtures of copper phthalocyanines and azopyridone yellow dyes (such as the dyes exemplified in JP2701387B2 [Mitsubishi Kasei Corp]) are unsatisfactory for use in colour filter applications because of poor light fastness. However, green colour filters containing azopyridone yellow dyes are significantly brighter than those produced using

alternative yellow chromophores such as azobenzenes, azopyrazolones, metallised azo dyes and yellow pigments, which are prevalent in the art.

It is desirable therefore to find improved colorants for use in colour filters, in particular colorants which can be used either alone for example as a yellow or as a yellow component in a mixture (e.g. with cyan colorants such as phthalocyanines). It is a preferred object of a preferred aspect of the invention to discover a green colorant comprising a mixture of one or more yellow colorants with one or more cyan colorants which is of particular use in a colour filter, optionally with improved properties such as light fastness and/or brightness.

The applicant has now found that a specific class of azopyridone colorants have utility in preparing patterned colored substrates such as colour filters. The specific class of colorants give stable inks with advantageous properties. As a preferred aspect of the present invention the applicant has surprisingly discovered certain simple pyridone dyes containing carboxy groups ortho to the azo linkage have a significant advantage in terms of light fastness versus related ortho-sulpho analogues, with no loss in brightness.

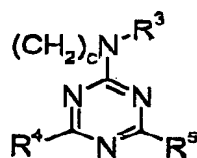
Therefore broadly in accordance with the present invention there is provided a compound of Formula 1



Formula 1

in which:

R¹ represents H, an optionally substituted C₁₋₃carbyl derived group, or a group of Formula A



Formula A

where:

c is from 2 to 6,

R^a represent H or optionally substituted C₁₋₃carbyl derived group; and

R⁴ and R⁵ independently represent an optional substituent;

R² represents an optionally substituted C₁₋₃carbyl derived group.

X, Y and Z independently represent H, an optional substituent or optionally substituted C₁₋₃carbyl derived group;

M represents H or a cation

m and n independently represent 0, 1 or 2;

with the provisos that:

at least one of R^1 , R^2 , X, Y or Z comprises a group of formula SO_3M or PO_3M_2 where M is

independently as represented herein; and

when n is 0; m is 1; X is a sulpho group para to the azo group; Z is H and R^2 is methyl then R^1 is other than ethyl.

The azopyridones of formula 1 are particular suitable for use in colour filters, giving very bright yellow films with good light fastness, and as components of ink-jet inks.

Conveniently compounds of formula 1 are not pyridinium pyridones

Preferred compounds of Formula 1 are those in which:

R^1 represents H, optionally substituted alkyl, a group of formula $[(CH_2)_{a_1}O][(CH_2)_{a_2}O] \dots [(CH_2)_{a_b}O]R^6$, or a group of formula A

R^2 represents Alkyl, CH_2SO_3M ,

X and Y independently represent SO_3M , CO_2M , PO_3M_2 , $SO_2NR^6R^7$, $CONR^6R^7$, CO_2R^6 , COR^6 , alkyl, alkoxy, NR^6COR^7 , halogen, NO_2 , NR^7R^6 ;

where R^6 and R^7 independently represent H, optionally substituted C_{1-6} alkyl, optionally substituted aryl; and R^6 represents R^6 or a substituted triazinyl group;

Z represents CN, $CONH_2$, H or CH_2SO_3M ;

R^3 represents H or optionally substituted C_{1-6} carbonyl derived group;

R^4 and R^5 independently represent halo, SO_3M , NR^6R^7 , OR^6 or SR^7 ;

M represents H, alkali metal ion, ammonium, or a quaternary ammonium cation (hereinafter QAC); more preferably M is Li^+ , Na^+ , K^+ or NH_4^+ ;

m, n is 0, 1 or 2;

$a_1 - a_b$ is each independently 2 or 3;

b is an integer from 1 to 10; and

c is an integer from 2 to 6;

with the proviso that at least one of R^1 to R^6 , X, Y or Z comprises a group of formula SO_3M or PO_3M_2 ; and the compound of formula 1 is other than a compound of formula II, III or IV as described herein.

Preferred QAC's are those containing C_{1-30} alkyl chains. More preferred QAC cations may be selected from one or more of the following: N,N-diethyl-N-dodecyl-N-benzylammonium; N,N-dimethyl-N-octadecyl-N-(dimethylbenzyl) ammonium; N,N-dimethyl-N,N-didodecyl ammonium; N,N-dimethyl-N,N-didodecyl ammonium; N,N,N-trimethyl-N-tetradecylammonium; N-benzyl-N,N-dimethyl-N-(C_{12-16} alkyl)ammonium; N-(dichlorobenzyl)-N,N-dimethyl-N-dodecylammonium; N-hexadecyl pyridinium; N-hexadecyl-N,N,N-trimethylammonium, dodecylpyridinium; N-benzyl-N-dodecyl-N,N-bis(hydroxyethyl)ammonium; N-dodecyl-N-benzyl-N,N-dimethylammonium; N-benzyl-

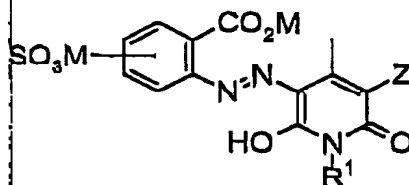
SMC 60384

5

N,N-dimethyl-N-(C₁₂₋₁₈ alkyl)ammonium; N-dodecyl-N,N-dimethyl-N-(1-naphthylmethyl) ammonium and N-hexadecyl-N,N-dimethyl-N-benzylammonium cations.

Suitable QAC cations may also be formed from suitable amines for example from one or more amines selected from: isononylamine, dodecylamine, octadecylamine, didecylamine, didodecylamine, tetradecylamine, hexadecylamine, mixed C₁₂₋₁₈ alkylamines and N-benzyl amines. Preferred amines which may be used to form suitable QAC comprise N-C₁₋₈ alkyl primary amines, N,N-di-C₁₋₈ alkyl secondary amines and N-benzyl amines. Particularly preferred amines comprise methyl and ethyl amine derivatives.

More preferred compounds of Formula 1 comprises dyes of formula B :



Formula B

in which

Z is CONH₂, CN or H

R¹ is optionally substituted C₂₋₈ alkyl (preferably hydroxy substituted) or a glycol group (for example, CH₂CH₂OCH₂CH₃ or CH₂CH₂OCH₂CH₂OH);

with the proviso that if the SO₃M group is in the 4 position of the benzene ring then either R¹ is other than ethyl or Z is other than H.

In general, preferred compounds of the invention are those which give particularly bright yellow films and prints, and are easily synthesised from readily available intermediates. Solubility in an aqueous ink is also desirable, hence compounds of formula 1 comprise at least one SO₃M or PO₃M₂ group, comprise substituent (e.g. R¹) of C₈ (or less) carbyl-derived groups. To improve solubility optionally R¹ comprises at least one PEG and/or OH group.

Compounds of the present invention can be prepared by analogy to any of the methods known in the art, for example GB 1,271,226 (ICI).

According to a further aspect of the present invention there is provided an ink comprising a solvent (preferably water) and at least one colorant of Formula (1):

When in the formulae herein there is a list of labels (e.g. Ar¹ and Ar²) or indices (e.g. 'n') which are said to represent a list of groups or numerical values, and these are said to be "independent in each case" this indicates each label and/or index can represent any of those groups listed: independently from each other, independently within each repeat unit, independently within each Formula and/or independently on each group which is substituted; as appropriate. Thus in each of these instances many different groups might be represented by a single label (e.g. Ar¹).

The term 'halo' as used herein signifies fluoro, chloro, bromo and iodo.

The terms 'optional substituent' and/or 'optionally substituted' as used herein (unless followed by a list of other substituents) signifies the one or more of following groups (or substitution by these groups): carboxy, sulpho, phospho, formyl, hydroxy, amino, imino, nitrilo, mercapto, cyano, nitro, halo and/or combinations thereof. These optional groups include all chemically possible combinations in the same moiety of a plurality (preferably two) of the aforementioned groups (e.g. amino and sulphonyl if directly attached to each other represent a sulphamoyl radical).

The term 'carbyl-derived' as used herein denotes any monovalent or multivalent organic radical moiety which comprises at least one carbon atom either solely (e.g. $-C\equiv C-$) or optionally combined with at least one other non-carbon atom (e.g. alkyl, carbonyl etc.). The non-carbon atom(s) may comprise any elements other than carbon (including any chemically possible mixtures or combinations thereof) that together with carbon can comprise an organic radical moiety. Preferably the non-carbon atom is selected from at least one hydrogen and/or heteroatom, more preferably from at least one: hydrogen, phosphorus, halo, nitrogen, oxygen and/or sulphur, most preferably from at least one hydrogen, nitrogen, oxygen and/or sulphur. Carbyl-derived groups include all chemically possible combinations in the same group of a plurality (preferably two) of the aforementioned carbon and/or non-carbon atom containing moieties (e.g. alkoxy and carbonyl if directly attached to each other represent an alkoxycarbonyl radical).

Preferably 'carbyl-derived' moieties comprise at least one of the following carbon containing moieties: alkyl, alkoxy, alkanoyl, carboxy, carbonyl, formyl and/or combinations thereof; optionally in combination with at least one of the following heteroatom containing moieties: oxy, thio, sulphinyl, sulphonyl, amino, imino, nitrilo and/or combinations thereof.

The term 'hydrocarbyl' as used herein (which is encompassed by the term 'carbyl-derived') denotes any radical moiety which consists only of at least one hydrogen atom and at least one carbon atom. A hydrocarbyl group may however be optionally substituted.

More preferably 'hydrocarbyl derived' moieties comprise one or more of the following carbon containing moieties: alkyl, aryl, alkaryl and/or combinations thereof. The term 'aryl' as used herein signifies a radical which comprises an aromatic hydrocarbon ring, for example phenyl, naphthyl, anthryl and phenanthryl radicals. The term 'alkyl' or its equivalent (e.g. 'alk') as used herein may be readily replaced, where appropriate, by terms denoting a different degree of saturation and/or valence e.g. moieties that comprise double bonds, triple bonds, and/or aromatic moieties (e.g. alkenyl, alkynyl and/or aryl) as well as multivalent species attached to two or more substituents (such as alkylene).

Any radical group mentioned herein as a substituent refers to a monovalent radical unless otherwise stated. A group which comprises a chain of three or more atoms signifies a group in which the chain may be straight or branched or the chain or part of the

chain may form a ring. For example, an alkyl group may comprise: propyl which includes n-propyl and isopropyl; butyl which includes n-butyl, sec-butyl, isobutyl and tert-butyl; and an alkyl group of three or more carbon atoms may comprise a cycloalkyl group. The total number of certain atoms is specified herein for certain substituents, for example C_{1-n} alkyl, signifies an alkyl group having from 1 to n carbon atoms. Preferred alkyl groups in dyes of Formulae 1 and 2 may be branched or straight chain and preferred branched chain alkyl groups are α -branched alkyl groups.

Advantageously the optional substituents and/or carbyl derived groups which may be present in Formula 1 herein may be each independently selected from: carboxy, sulphy, phospho, nitro, bromo, chloro fluoro, alkyl (especially C₁₋₄alkyl) alkoxy (especially C₁₋₄alkoxy), hydroxy, sulphyamoyl, amine (especially -NHR' and NR'Ar), mercapto, thioalkyl (especially C₁₋₄thioalkyl), cyano, ester (especially OCOR' or COOR') and amide (especially CONHR' and NHCOR');

where R' is H or optionally substituted C₁₋₄alkyl (especially H or C₁₋₄alkyl) and Ar is an optionally substituted aromatic ring (especially a benzene ring).

More preferred optional substituents and/or carbyl derived groups in Formula 1 are selected from SO₃M, CO₂M, PO₃M₂, Cl, Br, F, OH, C₁₋₄alkyl, C₁₋₄alkoxy, CONH₂, SO₂NH₂, OCO(C₁₋₄alkyl), COO(C₁₋₄alkyl), NHar and NHCO(C₁₋₄alkyl), where M and Ar are as defined herein.

Certain compounds and/or moieties therein (such as repeat units), which comprise the present invention may exist in many different forms for example at least one form from the following non-exhaustive list: salts (e.g. with organic and/or inorganic acids and/or bases including acid and/or base addition salts); isomers, stereoisomers, enantiomers, diastereoisomers, geometric isomers, tautomers, conformers, zwitterions, forms with regio-isomeric substitution, isotopically substituted forms, polymorphs, polymeric configurations, tactic forms, interstitial forms, complexes, chelates, clathrates, interstitial compounds, non-stoichiometric complexes, stoichiometric complexes, ligand complexes, organometallic complexes, solvates, isotopic forms, mixtures thereof and/or combinations thereof within the same species. The present invention preferably comprises all such forms of compounds, polymers, moieties therein, any compatible mixtures thereof and/or any combinations thereof, which comprise the present invention, preferably those which are effective in IJP and/or colour filters.

Salts of Formula 1 may be formed from one or more organic and/or inorganic bases. Preferred salts of Formula 1 are soluble in water.

The inks preferably contain from 1 to 10, more preferably from 1 to 6, especially from 1 to 3, more especially 1 dye of Formula 1.

The dyes of Formula 1 are preferably soluble in water. However, they may be modified to be soluble in organic solvents by use of a QAC as the counterion, where QAC is as hereinbefore defined.

The dyes of Formula 1 may be prepared by any suitable method known in the prior art.

The ink may contain further dyes other than those of Formula 1, for example to modify the colour or brightness of the ink.

5 The dye could be used individually as part of a YMC (yellow, magenta, cyan) colour filter, or mixed with another dye or dyes as the red or green component of an RGB (red, green, blue) colour filter. The arrangement of pixels could be any of those known in the art (stripe, mosaic, delta) and the filters would be suitable for displays, especially LCD's, and solid state imaging devices.

10 Use of the above dyes gives filters with very good fastness properties and brighter than those in the prior art.

The dyes may also be used in an ink-jet printing ink to give bright yellow prints with good fastness properties.

15 It is another object of the present invention to provide improved inks for processes for forming film coatings, that overcome some or all of the disadvantages of the prior art as discussed above, as well as products made and coated by processes using such inks (in particular colour filters).

20 Therefore according to another aspect of the present invention, there is provided an ink containing a dye of Formula 1 for use in any process for preparing a patterned, cross-linked, polymer, film coating on a substrate.

25 It is particularly preferred that the inks of the current invention are used in processes to manufacture a colour filter. These processes may comprise or consist of steps known in methods for producing colour filters (with colorants other than the novel colorants of the present invention). Such processes are well known in the art and include various printing, photolithographic, photographic, electrodeposition, laser ablation and thermal transfer processes. Examples of suitable processes are described below and in the following references, but it is to be understood that the invention is not limited to these processes :

30 "Reliability Improvements of Dichromated Gelatin Color Filters for TFT-LCD's", A. Endo, E. Hirose, T. Sato, S. Otera, N. Chiba, Polym. Mater. Sci. Eng., 1990, 63, 472-6.

"Process and Material for Color Filter Preparation in Liquid Crystal Display", H. Aruga, J. Photopolym. Sci. Technol., 1990, 3, 9-16.

"Color Filter for Liquid Crystal Display", S. Okazaki, Trans. Inst. Electron. Inf. Commun. Eng., Sect. E, 1988, E71, 1077-9.

35 P. Gregory, Chapter 2 "Micro Color Filters" in "High-Technology Applications of Organic Colorants", Kluwer Academic/Plenum Publishers, 1991.

"Color Filters for LCDs", K. Tsuda, Displays, 1993, 14, 115-24.

"Printing Color Filter for Active Matrix Liquid Crystal Display Color Filter", K. Mizuno, S. Okazaki, Jpn. J. Appl. Phys., Part 1, 1991, 30, 3313-17.

SMC 60384

9

EP 661350 (=US 5,608,091) (Nippon Shokubai)

EP 833203 (Nippon Shokubai)

It will be appreciated that where the colorants described are not water soluble dyes, the processes described may not be suitable for, or may need appropriate
5 modification to allow the use of, colorants of the present invention. Alternatively, where the colorants described are solvent soluble dyes, colorants of the present invention may be used by modification of their solubility in organic solvents through appropriate choice of the counter ion as described previously.

A typical process in which a dye of the present invention may be used comprises
10 the steps of:

(a) applying to the substrate simultaneously and/or sequentially in any order:

(i) one or more cross-linkable precursor(s) for one or more cross-linked polymer(s);

(ii) optionally one or more separate cross-linker(s) capable of cross-linking the precursor(s) for the polymer(s); and

15 (iii) one or more colorant(s) of Formula (1) optionally with one or more other colorant(s);

(b) optionally patterning one or more non cross-linked film(s) of component (i); component (ii); component (iii) and/or mixture(s) thereof, optionally before application of further components; and

20 (c) initiating cross-linking the mixture of components (i); (ii) in situ, to form an optionally patterned, cross-linked polymeric film coating on the substrate.

Preferably the application method in step (a) comprises applying an ink comprising both components (i) and (ii).

A process for which the present invention is particularly suitable is a printing
25 process, especially an ink-jet printing (IJP) process.

Preferably the printing process used is thermal or piezo IJP. The principles and procedures for ink jet printing are described in the literature for example in High Technology Applications of Organic Colorants, P. Gregory, Chapter 9 ISBN 0-306-43637-X.

30 Other suitable printing methods comprise: flexographic printing; off-set printing [e.g. as described in JP-A-10(98)-088055 (Sumitomo Rubber)] lithographic printing; gravure printing; intaglio printing; dye diffusion thermal transfer; screen and/or stencil printing [e.g. as described in WO 97-048117 (Philips Electronics)] and/or using 'typographic ink imaging pins' (e.g. as described in WO 97-002955 (Corning Inc.)).

35 Preferred methods of applying the polymer precursor and colorant of Formula (1) to the substrate comprise one or more of the following.

1) print onto the substrate (advantageously by IJP) a mixture comprising the polymer precursor which is thermally cross-linkable and the colorant; and thereafter curing the

mixture in situ (e.g. as described in the applicant's co-pending application GB 9824818.0).

2) Apply to the substrate a polymer precursor which is an anionic colourable photosensitive resin, and then exposing the resin to UV light via a mask to either make the exposed portions, which correspond to the pixels accept colorant; or harden the resin at the exposed portions, which correspond to the black matrix, to make it resistant to colorant; and print onto the resin (advantageously by IJP) a solution of the colorant. [e.g. as described in EP 0703471(Canon)]:

An additional method of applying the ink is by a photolithographic process. This may involve either:

1) Apply to a substrate a polymer precursor which is an anionic colourable photosensitive resin; and then expose the resin to radiation (e.g. UV light) through a patterned mask, develop the substrate to remove unexposed portions of the resin, optionally heat to further set the resin, then dip the substrate into an aqueous solution of the colorant. [e.g. as described in US 5,190,845 (Nippon Kayaku)]

2) Apply to a substrate an ink containing a photosensitive resin and the colorant; and then expose the resin to radiation (e.g. UV light) through a patterned mask, develop the substrate to remove either the exposed or unexposed portions of the resin and optionally heat to further set the resin [e.g. as described in EP 564237 (Mitsui Toatsu)]

3) Apply to a substrate a non-photosensitive ink containing the colorant and a thermally cross-linkable resin; then apply an ink containing a photosensitive resin over this coloured film; expose the photosensitive resin to radiation (e.g. UV light) through a patterned mask; develop the substrate to remove either the exposed or unexposed portions of the photosensitive resin and the corresponding portions of the coloured film beneath, heat to thermally cure the coloured polymeric film and optionally strip the photosensitive resin that remains [e.g. as described in US 5,176,971 (Kyodo Printing) and WO 88/05180 (Brewer Science Inc.)]

In (1) the photosensitive resin may be either a natural polymer such as gelatin or casein which has been photosensitised by the addition of for example ammonium dichromate, or may be a synthetic polymer.

In (2) and (3) the photosensitive resins used may be of either the positive or negative type. In the positive type, the solubility of the resin in a developing solution increases on exposure to radiation; in the negative type the solubility of the resin in a developing solution decreases on exposure to radiation.

In the laser ablation method, an ink containing the colorant and a (optionally thermally curable) non-photosensitive resin is applied to the substrate, then portions of the substrate are irradiated with a laser beam to selectively remove the ink in those areas through vaporisation and the remaining ink is optionally heated to thermally cure the resin [e.g. as described in JP10274709 (Sekisui Chem Ind.)]

SMC 60384

11

In these processes the inks may be applied to the substrate by any known coating method including spin-coating, bar-coating, dip-coating, curtain-coating, roller-coating and electrospray.

5 The process of the present invention can be used to give optionally patterned, optionally transparent films and coatings on substrates in general, including substrates which are not transparent. Accordingly the present invention includes a process for preparing polymeric film coatings for substrates in general not just colour filters.

10 The cross-linked polymeric film coating may be formed on a substrate to which the coating will bond, adhere, absorb or fuse. Preferably (e.g. if the process of the present invention is used to manufacture a colour filter) the substrate is transparent. Suitable transparent substrates include glass; plastics films and plates such as those of polyvinylalcohol, polyester, polyvinylchloride, polyvinylfluoride, polycarbonate, polystyrene, polyamide or polyimide. The substrate may be flexible or may be a flat panel (e.g. as used in many LCD displays). A preferred substrate is glass.

15 The substrates may be pre-treated to improve bonding, adhesion, absorption, fusion or spreading of the cross-linked polymeric coating on the substrate. Suitable pre-treatments include plasma etching in which the substrate is placed in an oxygen atmosphere and subjected to an electrical discharge or application of an adhesion promoter such as a silane.

20 An ink suitable for manufacture of a colour filter according to the present invention may be made by any method known in the art and comprise: one or more colorant(s) of Formula (1), one or more solvents and optionally other formulating agents. The inks may in addition contain precursor(s) for cross-linked polymer(s), one or more cross-linker(s) capable of cross-linking the precursor(s), optionally one or more non cross-linkable polymer(s) for improving the film-forming ability of the inks or the properties of the final films and (as appropriate for chemically or photochemically initiated systems) either a radical source, a photopolymerisation initiator or a dissolution inhibitor. An ink coloured in one of the desired colours can be produced with dyes of the present invention and optionally one or more other colorants, typically either yellow, green or red.

25 30 Preferably the optionally patterning method in step (b) of the process of the present invention uses electromagnetic radiation, more preferably UV radiation. Optionally to produce a colour filter the pattern formed may comprise of a multiplicity of discrete filter regions (pixels) on a transparent substrate via a single pass ink-jet printing process. Optionally, the transparent substrate has previously been subdivided into discrete pixel regions by any method known in the art (for example formation of a black matrix by photolithography).

35 The steps of the process of the present invention described herein may be followed for each of the desired colours to form a multi-colour optical filter structure so that the filter structure finally comprises the transparent substrate and a single layer of

differently coloured pixels arranged in triads or in any desired groups, each consisting of a predetermined number of differently coloured pixels.

The inks of the present invention are particularly useful for forming the green and red pixels of an additive (red, green and blue [RGB]) colour filter and for forming the yellow pixels of a subtractive (yellow, magenta and cyan [YMC]) colour filter.

A further aspect of the invention provides a process for printing an image on a substrate comprising applying thereto an ink according to another aspect of the present invention preferably by printing more preferably by means of an ink jet printer. Preferably the ink comprises solvent (preferably aqueous) and a dye of Formula (1) as described herein.

The ink jet printer preferably applies the ink to the substrate in the form of droplets which are ejected through a small nozzle onto the substrate. Preferred ink jet printers are piezoelectric ink jet printers and thermal ink jet printers. In thermal ink jet printers, programmed pulses of heat are applied to the ink in a reservoir (e.g. by means of a resistor adjacent to the nozzle) thereby causing the ink to be ejected in the form of small droplets directed towards the substrate during relative movement between the substrate and the nozzle. In piezoelectric ink jet printers the oscillation of a small crystal causes ejection of the ink from the nozzle. The term ink-jet printer denotes any device which could use an IJP technique to produce an image.

A further aspect of the present invention provides a substrate which has applied thereon an ink of the present invention as defined herein and/or which has been prepared by the process of the present invention also as defined herein.

The substrate, which is optionally transparent, preferably comprises plastic, metal, glass, paper, an overhead projector slide and/or a textile material. More preferably for a colour filter the substrate is glass. Preferred textile materials for ink jet printing are cotton, polyester and blends thereof. When the substrate is a textile material the process for printing an image thereon according to the invention preferably further comprises the step of heating the resultant printed textile, preferably to a temperature of 50°C to 250°C.

According to a further feature of this invention there is provided a colour filter comprising red, green and blue filter elements, or yellow, magenta and cyan filter elements, and comprising a coloured cross-linked polymeric coating on a transparent substrate prepared by the processes described in the present invention.

A further feature of the invention comprises a display containing a colour filter prepared according to the present invention.

A further feature of the present invention is a cartridge suitable for use in an ink jet printer containing an ink according to the invention. Also there is provided an ink jet printer containing an ink according to the invention.

SMC 60384

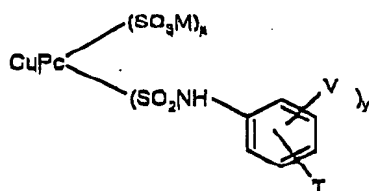
13

The colorants of the present invention may be particularly good yellow dyes which form particularly good green colorants when combined with a further cyan dye or pigment. Such compositions are particularly useful to produce a green colour filter

For the purposes of the present invention the term "colorant" as used herein denotes perceptible and/or emissive materials. The term "perceptible material" as used herein includes all dyes and/or pigments and denotes materials which absorb radiation substantially in that part of the electromagnetic (EM) spectrum which encompass the infra red (IR); visible and/or ultraviolet (UV) regions, preferably in a region where the radiation wavelength $[\lambda]$ is from about 200 nm to about 800 nm, more preferably in the visible region which is detectable by the normal, unaided human eye. The term "emissive material" as used herein denotes a material which is capable of emitting radiation, preferably EM radiation, more preferably radiation in the IR, visible and/or UV regions of the EM spectrum. Examples of emissive materials comprise fluorescent, phosphorescent and/or radioactive materials.

Therefore broadly in another aspect of the present invention there is provided a coloured composition comprising a compound of formula 1 or 2 (as described herein) together with one or more cyan or green dye(s) and/or one or more cyan or green pigments(s).

Preferably the cyan or green dye comprises a water soluble metal phthalocyanine, more preferably a copper, zinc, aluminium and/or nickel phthalocyanine. Most preferred cyan or green dyes comprises those of the following formula :



in which:

T independently represents the same as given for X or Y compounds of formula 1, preferably H, alkyl, alkoxy, CO_2M , SO_3M ;

V represents CO_2M , SO_3M or PO_3M_2 , especially meta- CO_2M

x and y independently represent from 0 to 4, preferably from 1 to 3; and

x + y is from 3 to 5, preferably 4;

where M is as given herein for compounds of Formula 1 herein.

Where the cyan or green colorant is a pigment, preferred pigments include C.I. Pigment Green 7 and C.I. Pigment Green 36.

The colorants of the present invention may also be used as shading components in combination with red, magenta and/or orange colorants to produce good red colorants which are useful for producing a red colour filter.

Therefore, broadly in another aspect of the present invention there is provided a coloured composition comprising a compound of Formula 1 (as described herein) together with one or more red, magenta and/or orange colorant(s) . [e.g. dye(s) and/or pigment(s)].

5 It is also to be understood that one or more further yellow colorant(s) , [e.g. dye(s) and/or pigment(s)]. other than those of the current invention, may be present in the ink for the purpose of adjusting the shade or enhancing the fastness properties of the ink and the resultant film or image produced using the ink.

10 Apart from the colorants of the present invention of formula 1 and 2, inks, colour filters and processes of the present invention may comprise further colorants to form a colorant mixture.

15 The further colorants (as well as colorants of the present invention) are preferably compatible with the resultant cross-linked polymer coatings, i.e. the resultant cured films have high transparency. Where the colorant is a dye, preferably the colorant is insoluble in organic solvents and soluble in water, for example the colorant may contain sulpho, phospho or carboxy groups.

20 If the colorant used is not soluble in the solvent used for the ink, the colorant is preferably present as a fine dispersion, prepared by for example milling the colorant in a solvent in a horizontal shaker in the presence of glass or metal beads and a dispersant. Suitable dispersants may comprise an anionic type (for example lignosulphonates and other sulphonated aromatic species) or a non-ionic type (for example alkylene oxide adducts).

25 Useful classes of further colorants include azos (including metallised azos), anthraquinones, phthalocyanines, perylenes, quinacridones, diketopyrrolopyrroles, pyrrolines, thiophenedioxides, triphenodioxazines, methines, benzofuranones, benzodifuranones, coumarins, indoanilines, benzenoids, xanthenes, triphenylmethanes, nitros, nitrosonaphthols, phenazines, solvent soluble sulphur dyes, quinophthalones, pyridones, aminopyrazoles, pyrrolidines, pyrroles, styrylics, maleimides, triphenazonaphthylamines, styryls, dithienes, azomethines, cyanines, indanthrones, benzimidazolones, isoindolinones, isoindolines and azoics.

30 The Colour Index International lists suitable dyes and pigments for use as further colorants such as acid dyes, direct dyes, basic dyes, reactive dyes, solvent dyes, disperse dyes and pigments and further examples of acid dyes are given in the Colour Index, 3rd Edition, Volume 1, pages 1003 to 1561, further examples of direct dyes are given in Volume 2, pages 2005 to 2478, further examples of basic dyes are given in Volume 1, pages 1611 to 1688, further examples of reactive dyes are given in Volume 3 pages 3391 to 3560, further examples of solvent dyes are given in Volume 3, pages 3563 to 3648, further examples of disperse dyes are given in Volume 2, pages 2479 to 2742,

SMC 60384

15

and further examples of pigments are given in Volume 3 pages 3267 to 3390,. These colorants are included herein by reference.

Subject to the provisos herein, generally preferred colorants are pigments; or dyes which have substituent groups which aid the solubility of the dye(s) in liquid media used in the process or which aid the solubility of the dye(s) in the final cross-linked polymer matrix.

Preferably in one aspect of the present invention there comprises an ink effective for IJP, the ink comprising:

(a) from 0.01 to 30 parts of an IJP-effective compound of Formula (1); and
(b) from 70 to 99.99 parts of a liquid medium or a low melting point solid medium; wherein all parts are by weight and the number of parts of (a) + (b) = 100.

The number of parts of component (a) is preferably from 0.1 to 20, more preferably from 0.5 to 15, and especially from 1 to 5 parts. The number of parts of component (b) is preferably from 99.9 to 80, more preferably from 99.5 to 85, especially from 99 to 95 parts.

When the medium is a liquid, preferably component (a) is completely dissolved in component (b). Preferably component (a) has a solubility in component (b) at 20°C of at least 10%. This allows the preparation of concentrates which may be used to prepare more dilute inks and reduces the chance of the colorant precipitating if evaporation of the liquid medium occurs during storage.

Preferred liquid media include water, a mixture of water and a water miscible organic solvent.

When the medium comprises a mixture of water and an organic solvent, the weight ratio of water to organic solvent is preferably from 99:1 to 1:99, more preferably from 99:1 to 50:50 and especially from 95:5 to 80:20. The liquid medium may comprise water and preferably two or more, more preferably from 2 to 8, water-soluble organic solvents.

The water-miscible organic solvent(s) may comprise any of the following and/or mixtures thereof: C₁₋₆-alkanols, preferably methanol, ethanol, n-propanol, isopropanol, n-butanol, sec-butanol, tert-butanol, n-pentanol, cyclopentanol and/or cyclohexanol; amides, preferably linear amides, for example dimethylformamide and/or dimethylacetamide; ketones and/or ketone-alcohols, preferably acetone, methyl ether ketone, cyclohexanone and/or diacetone alcohol; water-miscible ethers, preferably C₂₋₄ethers, tetrahydrofuran and/or dioxane; diols, preferably alkylene glycols containing a C₂-C₆ alkylene group; more preferably C₂₋₁₂diols (for example pentane-1,5-diol, ethylene glycol, propylene glycol, butylene glycol, pentylene glycol, hexylene glycol); thioglycols preferably thiodiglycol; oligo- and/or poly-alkyleneglycols (for example diethylene glycol, triethylene glycol, polyethylene glycol and/or polypropylene glycol); triols, preferably glycerol and/or 1,2,6-hexanetriol; lower alkyl glycol and polyglycol ethers, preferably C₁₋₄alkyl ethers of diols or mono-C₁₋₄alkyl ethers of C₂₋₁₂diols; (for example 2-methoxyethanol, 2-(2-methoxyethoxy)ethanol, 2-(2-ethoxyethoxy)ethanol, 2-(2-butoxyethoxy)ethanol, 3-butoxypropan-1-ol,

2-[2-(2-methoxyethoxy)ethoxy]ethanol, 2-[2-(2-ethoxyethoxy)ethoxy]ethanol and/or ethyleneglycol monoallyl ether); cyclic amides, preferably optionally substituted pyrrolidones (for example 2-pyrrolidone, N-methyl-2-pyrrolidone, N-ethyl-2-pyrrolidone, caprolactam and/or 1,3-dimethylimidazolidone); cyclic esters, preferably caprolactone; sulphoxides, preferably dimethyl sulphoxide and/or sulfolane.

More preferred water-soluble organic solvents are selected from: cyclic amides (for example 2-pyrrolidone, dimethyl pyrrolidone, N-methyl-2-pyrrolidone, N-ethyl-2-pyrrolidone N-(2-hydroxyethyl)-2-pyrrolidone and mixtures thereof.); diols, (for example 1,5-pentane diol, ethyleneglycol, thiodiglycol, diethyleneglycol and triethyleneglycol); C_{1-8} alkyl ethers of diols (for example 2-methoxy-2-ethoxy-2-ethoxyethanol); C_{1-8} -alkyl mono ethers of C_{2-8} -alkylene glycols; C_{1-8} -alkyl mono ethers of poly(C_{2-8} -alkylene glycols); and mixtures thereof.

A preferred liquid medium comprises:

- (a) from 75 to 95 parts water; and
- (b) from 25 to 5 parts in total of one or more solvents selected from: diethylene glycol, 2-pyrrolidone, thiodiglycol, N-methylpyrrolidone, cyclohexanol, caprolactone, caprolactam and pentane-1,5-diol;

where the parts are by weight and the sum of the parts (a) + (b) = 100.

Another preferred liquid medium comprises:

- (a) from 60 to 80 parts water;
- (b) from 2 to 20 parts diethylene glycol; and
- (c) from 0.5 to 20 parts in total of one or more solvents selected from: 2-pyrrolidone, N-methylpyrrolidone, cyclohexanol, caprolactone, caprolactam, pentane-1,5-diol and thiodiglycol;

where the parts are by weight and the sum of the parts (a) + (b) + (c) = 100.

Examples of further IJP-effective media for inks of the present invention comprise a mixture of water and one or more organic solvents are described in US 4,963,189, US 4,703,113, US 4,626,284 and EP 0425150-A.

When the liquid medium comprises an organic solvent free from water, (i.e. less than 1% water by weight) the solvent preferably has a boiling point of from 30° to 200°C, more preferably of from 40° to 150°C, especially from 50 to 125°C. The organic solvent may be water-immiscible, water-miscible or a mixture of such solvents. Preferred water-miscible organic solvents comprise any of those described above and mixtures thereof. Preferred water-immiscible solvents comprise aliphatic hydrocarbons; esters (for example ethyl acetate) chlorinated hydrocarbons (for example dichloromethane), ethers (for example diethyl ether) and mixtures thereof.

When the liquid medium comprises a water-immiscible organic solvent, preferably it comprises a polar solvent (for example a C_{1-4} alkanol) to enhance the solubility of the dye in the liquid medium. It is especially preferred that where the liquid medium is an

SMC 60384

17

organic solvent free from water it comprises a ketone (especially methyl ethyl ketone) and/or an alcohol (especially a C_{1-4} alkanol, more especially ethanol or propanol).

The organic solvent free from water may be a single organic solvent or a mixture of two or more organic solvents. It is preferred that when the medium is an organic solvent free from water it is a mixture of 2 to 5 different organic solvents. This allows a medium to be selected which gives good control over the drying characteristics and storage stability of the ink.

Ink media comprising an organic solvent free from water are particularly useful where fast drying times are required and particularly when printing onto hydrophobic and non-absorbent substrates, for example plastics, metal and glass.

Preferred low melting solid media have a melting point in the range from 60°C to 125°C. Suitable low melting point solids include long chain fatty acids or alcohols, preferably those with C_{18-24} chains, and sulphonamides. The dye of Formula (1) may be dissolved in the low melting point solid or may be finely dispersed in it.

The ink may also contain additional components conventionally used in inks for IJP, for example viscosity and surface tension modifiers, corrosion inhibitors, biocides, fogging reducing additives and surfactants which may be ionic or non-ionic.

In a further aspect of the present invention there is provided a general purpose ink optionally for use in preparing a colour filter, the ink comprising a fluid medium, and one or more colorant(s) of Formula 1 herein. The precursor(s), cross-linker(s) and colorant(s) are as defined previously.

Preferably inks according to the invention are prepared by mixing together (i) a solution of a dye(s) and optionally water.

The amount of dye and solvent contained in the ink will vary according to the depth of shade required. Typically, however, the ink will comprise

(a) from 0.5 to 15 parts, more preferably 0.8 to 10 parts, especially 1 to 8 parts in total of one or more dyes of Formula (1);

(b) from 0 to 90 parts, more preferably from 50 to 80 parts of water; and

(c) from 0 to 90 parts, more preferably 0 to 60 parts of one or more water miscible organic solvent(s); and

(d) other ingredients from 0 to 50 parts, more preferably 0 to 30;

where all parts are by weight and the total number of parts of (a) + (b) + (c) + (d) add up to 100.

The water-miscible solvent may be one or more of those described herein, preferably with a solubility in water at 20°C of more than 50g/l.

Instead of parts (b) and (c) [the water and water-miscible organic solvent(s)] the ink may comprise one or more water-immiscible organic solvent(s).

Suitable water-immiscible organic solvents include aromatic hydrocarbons, e.g. toluene, xylene, naphthalene, tetrahydronaphthalene and methyl naphthalene; chlorinated

aromatic hydrocarbons, e.g. chlorobenzene, fluorobenzene, chloronaphthalene and bromonaphthalene; esters, e.g. butyl acetate, ethyl acetate, methyl benzoate, ethyl benzoate, benzyl benzoate, butyl benzoate, phenylethyl acetate, butyl lactate, benzyl lactate, diethyleneglycol dipropionate, dimethyl phthalate, diethyl phthalate, dibutyl phthalate, di (2-ethylhexyl) phthalate; alcohols having six or more carbon atoms, e.g. hexanol, octanol, benzyl alcohol, phenyl ethanol, phenoxy ethanol, phenoxy propanol and phenoxy butanol; ethers having at least 5 carbon atoms, preferably C₅₋₁₄ ethers, e.g. anisole and phenetole; nitrocellulose, cellulose ether, cellulose acetate; low odour petroleum distillates; turpentine; white spirits; naphtha; isopropylbiphenyl; terpene; vegetable oil; mineral oil; essential oil; and natural oil; and mixtures of any two or more thereof

The water-immiscible solvent preferably has a solubility in water at 20°C of up to 50g/l.

A further feature of the invention provides a composition comprising a water-dissipatable polymer and a dye of Formula (1). In these compositions the preferred water-dissipatable polymers and dyes are as described in following published patent application (Avecia case reference in brackets): WO95/34204 (SMC 40116X) and the following of the applicant's co-pending patent applications (Avecia case reference in brackets): PCT/GB99/00785 (SMC 60286); PCT/GB99/00784 (SMC 60287); PCT/GB99/00783 (SMC 60288); GB 9824818.0 (SMC 60324); GB 9827690.0 (SMC 60330); GB 9827894.8 (SMC 60331); and GB 9914446.1 (SMC 60365). These applications are hereby incorporated by reference.

Such compositions may be dissipated in water and optionally mixed with further ingredients to give an ink, for example with one or more organic solvents.

The other ingredients may comprise one or more formulating agents conventionally used in inks for example to improve the solubility of colorant in the ink and/or to improve the flow and handling properties of the ink. Thus for example the ink may comprise one or more humectant(s); rheological agent(s) [such as viscosity modifier(s) and/or surface tension modifier(s), for example wax(es) (e.g. beeswax) and/or clay(s) (e.g. bentonite)]; corrosion inhibitor(s), biocides (such as those available commercially from Avecia Limited under the trade name Proxel GXL or from Rohm and Haas under the trade name Kathon); fungicide(s); kogation reducing additives; IR absorber(s) (such as that available commercially from Avecia Limited under the trade name Projet 900NP); fluorescent brightener(s), (such as C.I. Fluorescent Brightener 179); and surfactant(s) (which may be ionic or non-ionic and include surface active agent(s) wetting agent(s) and/or emulsifier(s) such as those described in McCutcheon's Emulsifiers and Detergents 1996 International Edition or in Surfactants Europa 3rd Edition 1996 each of which is incorporated herein by reference).

SMC 60384

19

The ink may also comprise radical scavengers and/or UV absorbers to help improve light and heat fastness of the ink and resultant colour filter. Examples of such additives include: 2-hydroxy-4-methoxy-5-sulfobenzophenone; hydroxy phenylbenzotriazole; 4-hydroxy-TEMPO and transition metal complexes (such as nickel complexes of thiocarbamic acids). These additives are used typically in an amount from 30% to 60% by weight of the colorant, and are further described in "The Effect of Additives on the Photostability of Dyed Polymers", *Dyes and Pigments*, 1997, 33(3), 173-196 and JP-A- 04-240603 (Nippon Kayaku).

For an aqueous ink, the ink preferably has a pH from 3 to 12, more preferably from 4 to 11. The pH selected will depend to some extent on the desired cation for the colorant and the method of application. The desired pH may be obtained by the addition of a pH adjusting agent such as an acid, base or pH buffer. The amount of pH adjuster used will vary according to the desired pH of the ink, but typically a base may be present in an amount of up to 30 %. Where a liquid(s) is added to the mixture the printed substrate may be dried by heating or by air drying at ambient temperature to evaporate the liquid before the coating is cured or during curing.

Examples of suitable formulations of polymer precursor and colorant which may be used in the method of the present invention are also given in the applicant's co-pending applications GB 9824818.0 and EP 0764290, where it will be appreciated that the colorants of Formula (1) herein replace some or all of the colorants used in these prior art formulations.

Preferably the ink of the present invention comprises from about 10 to about 99.6, preferably from about 30 to about 99.5, more preferably from about 50 to about 99, parts of the fluid medium; and from about 90 to about 0.4 parts, preferably from about 70 to about 0.5, more preferably from about 50 to about 1, parts of the other ingredients; where all parts are by weight and the number of parts totals 100.

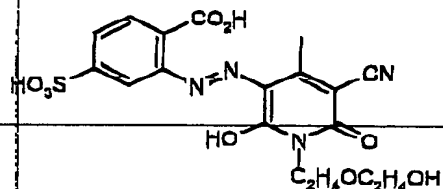
The inks according to a further aspect of the invention may be prepared by mixing the ingredients in any order. Suitable mixing techniques are well known in the art, for example agitation, ultrasonication or stirring of the components. The ingredients may be present in the ink in any form suitable for application to the substrate, for example the form of a dispersion, emulsification, suspension, solution or a combination thereof.

Examples of further media for inks of the present invention comprising a mixture of water and one or more organic solvents are described in US 4,963,189, US 4,703,113, US 4,626,284 and EP 0425150-A.

Dyes of the invention will now be illustrated by the following examples in which all parts and percentages are by weight unless specified otherwise. In the examples (and previously), compounds referred to by reference to CI numbers are the dyestuffs identified by these numbers in the Colour Index International, 3rd Edition, 3rd Revision. In

each of the following Examples, the inks were tested and the results were set out in tables.

Example 1



5- (2'-carboxy-5'sulphophenylazo)-1-(2'-hydroxyethoxyethyl)-3-cyano-4-methyl-6-hydroxy-2-pyridone

1(a) Preparation of coupler = 1-(2'-hydroxyethoxyethyl)-3-cyano-4-methyl-6-hydroxy-2-pyridone

Ethylcyanoacetate (115 parts) was slowly added to 2-(2-Aminoethoxy)ethanol (134 parts) with stirring, while maintaining the temperature below 30°C. On completion of the addition, the mixture was heated at 95°C for 3 hours then cooled back to room temperature. Methyl acetoacetate (123 parts) was added, maintaining the temperature below 30°C, the whole mixture then cooled to <10°C before adding ethylamine (57.6 parts). The resultant mixture was heated at 90°C for 18 hours, then cooled to room temperature. On acidification to pH1 with concentrated hydrochloric acid, the product precipitated out. This was isolated by filtering, washing with a small amount of 2N HCl and finally drying at 60°C to yield 81 parts of the above pyridone coupler.

1(b) Preparation of title compound:

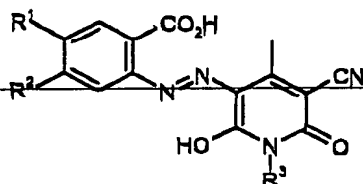
2-Amino-4-sulphobenzoic acid (27.2 parts) was stirred in water (200 parts) at pH5 until dissolved. The solution was cooled to <10°C and concentrated HCl (47 parts) was added. Sodium nitrite (6.8 parts) was slowly added and the reaction mixture was stirred at 0-10°C for 30 minutes. Excess nitrous acid was then removed by addition of sulphamic acid, until the diazo mixture was negative to starch-iodide. To this was then added a solution of the above pyridone coupler (53 parts) in water (200 parts), which had been adjusted to pH8 with 2N NaOH. The temperature of the reaction mixture was maintained below 10°C during the addition. The mixture was then stirred for 16 hours whilst warming to room temperature. The dark yellow solution was acidified to pH 1 with concentrated HCl and the resultant precipitate was collected by filtration. The precipitate was re-dissolved in water (200 parts), and adjusted to pH9 with 0.88 S.G. ammonia. The product was then re-precipitated by pouring on to concentrated HCl (35 parts) and collected by filtration. The product was dissolved in ammonia and re-precipitated and collected by filtration once more. Finally the product was dissolved in water (200 parts) containing ammonia at pH9,

SMC 60384

21

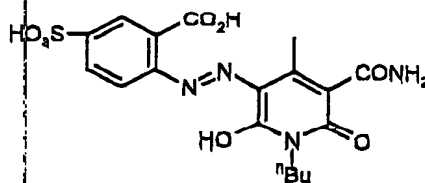
dialysed to remove the inorganic components, screened through a 0.45 μ filter and dried at 60°C to yield the title compound as a yellow powder of its ammonium salt (7.7 parts).

The following dyes were made by an analogous method to that described in Example 1:



Ex	R ¹	R ²	R ³
2	H	SO ₃ H	ⁿ Bu
3	SO ₃ H	H	ⁿ Bu
4	H	SO ₃ H	2-Ethylhexyl
5	SO ₃ H	H	2-Ethylhexyl
6	H	SO ₃ H	C ₂ H ₄ OH
7	H	SO ₃ H	C ₂ H ₄ NH ₂
8	H	SO ₃ H	CH ₂ C(CH ₃) ₃
9	H	SO ₃ H	C ₂ H ₄ OEt
10	H	SO ₃ H	Cyclohexyl
11	H	SO ₃ H	n-Hexyl
12	H	SO ₃ H	C ₈ H ₁₇ OEt
13	H	SO ₃ H	C ₈ H ₁₇ OH
14	SO ₃ H	H	C ₂ H ₄ OC ₂ H ₄ OH
15	SO ₃ H	H	C ₃ H ₇ OH

Example 16



5-(2'carboxy-4'sulphophenylazo)-1-(ⁿbutyl)-3-carbonamido-4-methyl-6-hydroxy-2-pyridone (S178363)

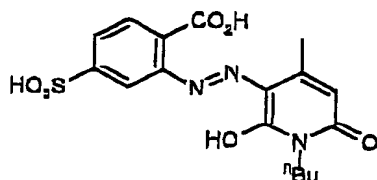
16(a) Preparation of coupler = 1-ⁿButyl-3-carbonamido-4-methyl-6-hydroxy-2-pyridone
1-ⁿButyl-3-cyano-4-methyl-6-hydroxy-2-pyridone (103 parts) was added carefully to 1.84 S.G. sulphuric acid (370 parts), maintaining the temperature below 50°C. After the

addition was complete the temperature was raised to 75°C for 12 hours. The reaction mixture was allowed to cool to room temperature, then drowned out onto ice (400 parts). The solution was carefully neutralised with calcium carbonate and screened; the solids were further washed through with copious amounts of water and ethanol. Finally, the solution was evaporated to dryness under reduced pressure and the solids dried at 60°C to give the above pyridone coupler as a light grey powder (122 parts).

16(b) Preparation of title compound

The title compound was made in an analogous manner to the dyestuff in Example 1, substituting a molar equivalent of the above pyridone coupler [prepared as described in 16(a)] for the coupler used in example 1(b), and a molar equivalent of 2-amino-5-sulphobenzoic acid for the diazo component used in example 1 (b).

Example 17



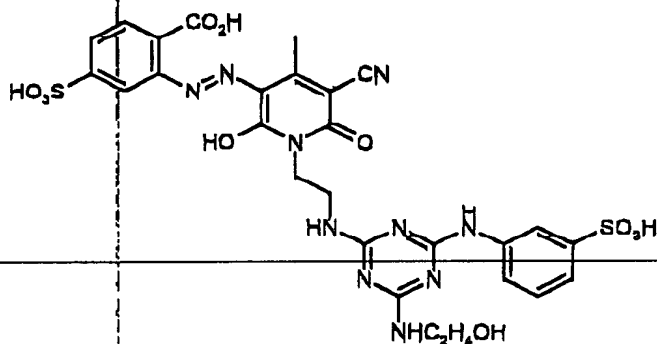
5-(2'carboxy-5'sulphophenylazo)-1-(ⁿbutyl)-4-methyl-6-hydroxy-2-pyridone

17(a) Preparation of pyridone coupler = 1-(ⁿButyl)-4-methyl-6-hydroxy-2-pyridone

1-(ⁿButyl)-3-cyano-4-methyl-6-hydroxy-2-pyridone (20.6 parts) was added slowly to 75% sulphuric acid (40.6 parts). After addition the solution was heated at 125°C for 3 hours then cooled back to room temperature. The thick oil was poured onto ice (200 parts) with good stirring. 47% Sodium hydroxide solution was then carefully added until the product started to solidify. After stirring for a further 30 minutes, the solid was collected by filtration and dried at 60°C to yield the title pyridone as a brown powder (17.5 parts).

17(b) Preparation of title compound

The title compound was made in an analogous manner to the dyestuff in Example 1, substituting a molar equivalent of the above pyridone coupler [prepared as described in 17(a)] for the coupler used in example 1(b).

Example 18

5 18(a) Preparation of coupler = 1-(2'-Aminoethyl)-3-cyano-4-methyl-6-hydroxy-2-pyridone
This coupler was made in an analogous manner to that described in Example 1(a), substituting ethylene diamine (5 times molar excess) for the 2-(2-aminoethoxy)ethanol.

10 18(b) Preparation of intermediate dye 5-(2'-carboxy-5'-sulphophenylazo)-1-(2'-aminoethyl)-3-cyano-4-methyl-6-hydroxy-2-pyridone

This dye was made in an analogous manner to the dyestuff in Example 1, substituting a molar equivalent of the above pyridone coupler [prepared as described in 18(a)] for the coupler used in example 1(b).

15 18(c) Preparation of title compound

Cyanuric chloride (2.6 parts) was dissolved in acetone (20 parts) and poured onto ice/water (100 parts) at 0-5°C. A solution of metanilic acid (2.4 parts) in water (50 parts) at pH7 was then added to the cyanuric chloride suspension, maintaining the reaction at 0-5°C and pH6. The mixture was stirred under these conditions for 2 hours. To this was then added a solution of 5-(2'-carboxy-5'-sulphophenylazo)-1-(2'-aminoethyl)-3-cyano-4-methyl-6-hydroxy-2-pyridone [5.2 parts, made as described in 18(b) above] in water (100 parts) at pH8.5, and the solution was stirred at pH8.5, 45°C for 12 hours. After cooling to room temperature the solution was acidified to pH 1 with concentrated HCl, and the solid was collected by filtration, dialysed and dried. This intermediate (8.7 parts) was then re-dissolved in water (400 parts) and ethanolamine (4.6 parts) was added. The solution was then heated at 70°C for 6 hours. After cooling to room temperature, the product was precipitated by acidifying to pH 1 with concentrated HCl and was collected by filtration. It was then re-dissolved in water (200 parts), adjusted to pH9 with 0.88S.G. ammonia and re-precipitated by pouring on to concentrated HCl (35 parts). After stirring for 10 minutes the product was collected by filtration then re-dissolved in ammonia and re-precipitated with concentrated HCl once more. After collection by filtration, the product was dialysed,

screened through a 0.45 μ filter and dried at 60°C to yield the ammonium salt of the title dye as a yellow powder (3.3 parts).

For the purpose of evaluating the spectra and resistance properties of formulations containing the dyes, the inks were bar-coated on to microscope slides using a No.1 K-bar (RK Print-Coat Instruments Ltd.), and the slides cured at 200°C for 15 minutes. The resultant films had a thickness of approximately 2 μ m.

Example 19

Inks with the following formulation were made (all parts by weight) :

Acrylic co-polymer (54% methyl methacrylate; 46% methacrylic acid)	13.2 parts
Primid XL552 - ex EMS Chemie	4.4 parts
2-Amino-2-methyl-1-propanol	13.2 parts
Dye	3.0 parts
Sodium dodecylbenzenesulphonate	0.9 parts
Water	65.3 parts

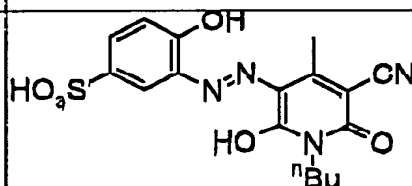
The inks gave very bright yellow films with high transmission at 520nm and low transmission at 420nm. The results are tabulated below, together with light fastness data

Ink Ex	Dye Ex	%T @ 420nm	%T @ 520nm	LF - ΔE /100 hours ¹
19	1	2	94	24.8
20	2	0	93	20.1
21	3	1	94	44.2
22	4	4	93	45.8
23	6	1	90	42.6
24	9	0	90	28.5
25	11	0	93	45.3
26	12	1	95	30.2
27	13	4	96	30.0
28	14	0	89	41.2
29	15	1	95	36.1
30	16	0	97	35.2
31	17	0	98	43.8
32	18	7	87	28.6
Comp. 1	A ²	14	65	77.3
Comp. 2	B ³	4	95	57.6

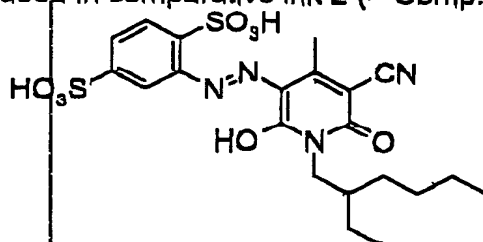
Footnotes

1. Light fastness properties were measured using a xenon lamp in an Atlas weatherometer Ci35A (lamp power 0.80 Wm^{-2} at 420nm, black panel 63°C , wet bulb depression 16°C), and the ΔE values determined using a Minolta CM-3600d spectrophotometer.

2. Structure of dye A used in comparative ink 1 (= Comp. 1)



3. Structure of dye B used in comparative ink 2 (= Comp. 2)

Example 33

Inks with the following formulation were made (all parts by weight) :

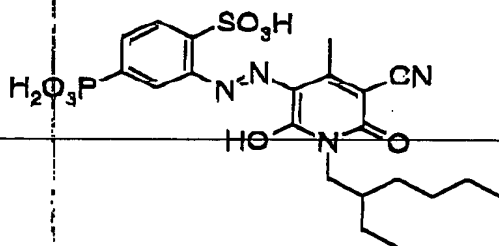
Acrylic co-polymer (34% methyl methacrylate; 46% methacrylic acid 20% hydroxyethyl methacrylate)	12.3 parts
Primid XL552 - ex EMS Chemie	4.1 parts
0.88 S.G. ammonia	12.3 parts
C.I. Reactive Blue 14	1.9 parts
Pyridone dye	3.3 parts
Sodium dodecylbenzenesulphonate	0.4 parts
Triethyleneglycol monobutyl ether	0.9 parts
Water	64.8 parts

The inks produced bright green highly transparent films, but the one made using a dye of this invention had far superior light fastness ;

Example	Pyridone dye	LF = $\Delta E/100$ hrs
33	4	10.6
Comp. 31	C	62.4

Footnotes

1. Structure of dye C used in comparative ink 3 (= Comp. 3):

Examples 34 to 39

Inks similar to that used in Example 33 were made, using an alternative cyan component (all parts by weight) :

Acrylic co-polymer (34% methylmethacrylate; 46% methacrylic acid 20% hydroxyethyl methacrylate)	11.5 parts
Primid XL552 - ex EMS Chemie	3.9 parts
0.88 S.G. ammonia	11.5 parts
Cyan D ¹	See Table
Pyridone dye of the present invention	See Table
Sodium dodecylbenzenesulphonate	0.4 parts
Triethyleneglycol monobutyl ether	2.9 parts
Water	to 100 parts

All the inks produced bright green highly transparent films, but the ones made using dyes of this invention had far superior light fastness :

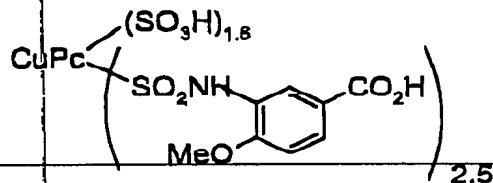
Example	Dye (%w/w)	Cyan D (% w/w)	Y ⁴	x	y	%T @ 540nm	LF = ΔE /100 hrs
34	1 (2.3)	2.5	59.7	0.315	0.557	84	13.8
35	2 (2.5)	3.0	59.1	0.311	0.560	83	16.9
36	9 (2.0)	2.5	59.1	0.306	0.550	83	15.1
37	16 (3.0)	2.0	55.0	0.287	0.578	80	13.3
38	17 (4.5)	1.5	55.2	0.301	0.571	78	14.0
39	E ² (2.9)	2.0	51.0	0.319	0.570	76	11.2
Comp. 4	F ³ (4.5)	1.5	58.6	0.298	0.560	83	42.3

SMC 60384

27

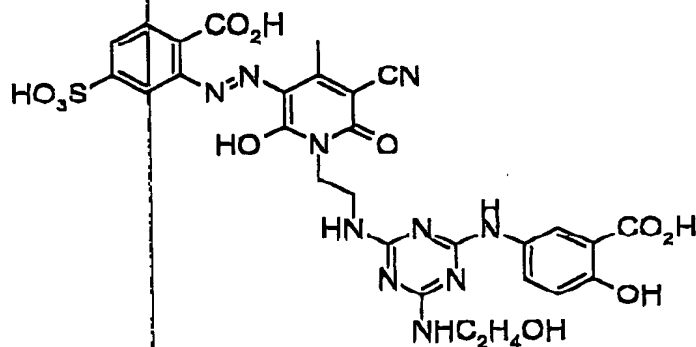
Footnotes

1. Structure of cyan dye D:

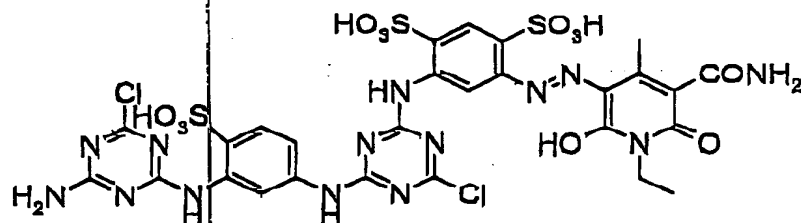


5

2. Structure of dye E used in Example 39 of the present invention:



3. Structure of dye F used in comparative ink 4 (= Comp. 4):



4. Chromaticity values were measured on a Minolta CM-3600d spectrophotometer using a C light source and 2° observer.

Example 40

A photocurable ink was produced by mixing together NeoRad R-441 (50 parts; 33% solids UV-curable resin commercially available from AVECIA Ltd.), NeoCryl BT-175 (5 parts; 40% solids acrylic resin commercially available from AVECIA Ltd.), 0.88 S.G. ammonia (2 parts), CGI 1700 (1 part; photoinitiator commercially available from Ciba), azopyridone dye of Example 3 (5 parts) and water (37 parts). The ink was applied to a glass substrate by bar-coating and exposed to UV-light through a photomask. The

unexposed portions of the film were then removed using a 1% sodium carbonate solution at 25°C and the remaining resin thermally cured at 150°C for 30 minutes to produce a bright yellow patterned film.

5 Example 41

To an ink base comprising 87.5 parts water, 5 parts 2-pyrrolidone, 5 parts thiodiglycol and 2.5 parts Surfynol 465™ (available from Air Products Ltd.) was added the dye of Example 2 to give a 4% solution of the dye in the ink base. The solution was then adjusted to pH 10 using NH₃ and filtered through a 0.45μ membrane.

10 When this was applied to paper using an ink-jet printer, strong bright yellow prints were obtained with excellent fastness properties.

Further inks

15 Further inks which are suitable for making colour filters may be prepared having the formulations described in tables below where the numbers denote parts by weight of each ingredient in the formulation. The dyes used may be in their free acid form and/or in the form of any IJP-effective salt. The number in the column headed Ex is the example number of a dye of the invention (1 to 18 and E) described herein with the number in brackets being the amount of that dye (w/w). The following abbreviations are also used in the tables:

20 DB199 = Direct Blue 199 DB86 = Direct Blue 86 DB87 = Direct Blue 87
 AB9 = Acid Blue 9 RB15 = Reactive Blue 15 CD = cyan D as given herein
 PAA = Poly(acrylic acid) of M_w 2000 X = Primid XL552
 TEA = Triethanolamine TMP = Trimethylolpropane W = Water
 25 NMP = N-methyl-2-pyrrolidone 2P = 2-pyrrolidone
 BE = Butoxyethanol DEG = Diethylene glycol GBL = γ-butyrolactone
 ADBS = Ammonium dodecylbenzene sulphonate
 SDBS = Sodium dodecylbenzene sulphonate
 SURF = Surfynol 465 (Non-ionic surfactant available from Air Products)
 30 AMP = 2-Amino-2-methyl-1-propanol AM = Ammonia (0.88 S.G.)
 HT = 4-Hydroxy-TEMPO; and
 HMBS = 2-Hydroxy-4-methoxy-5-sulfobenzophenone.

SMC 60384

29

TABLE I

Ex	DB199	DB86	PAA	X	W	BE	ADBS	AMP	AM
1(0.5)	3		15	5.5	60		1	15	
2(0.25)		3.	12	4	69.25	5	0.5		6
3(1)	3		12	6	62	5	1		10
4(0.2)	2		4	1	88.8		1	3	
5(0.5)		4	10	2	74.5	2			7
6(0.3)	3.5		12.5	4	61.4	5	0.8	12.5	
7(1)		1.5	8	2	80.5	2		5	
8(0.1)	2	0.9	11.5	6.5	63	5			11
9(3)		0.5	13.7	4.3	63.8		0.7	14	
10 (0.2)	2.8		15	5	61		1		15

TABLE II

Ex	DB199	PAA	X	TEA	TMP	W	2P	ADBS	AM
11(0.5)	4	14	5			61.5		1	14
12(0.5)	4	10.5	2.5			70	4	0.5	8
13(0.25)	3.5	15		5		60.5		0.75	15
14(0.1)	2.9	12			3	67	5		10
15(2)	0.5	12	2		2	76		0.5	5
16(0.1)	0.9	8	5			81	1		4
17(0.1)	3	15		5		63		0.9	13

TABLE III

Ex	DB199	PAA	TMP	W	2P	DEG	GBL	SURF	SDBS	AMP
18(0.5)	4	16	4	54.5			5	1		15
E(2.9)	0.1	12	3	71.2					0.8	10
1(3)	2	14	7	49	5	5				15
2(0.1)	1	4	1	87.4	0.5		1.5	0.5		4
3(0.5)	1.5	8	3	80.5		2			0.5	4

TABLE IV

Ex	AB9	CD	PAA	X	W	DEG	SURF	AMP	AM
4(3)	2		10	5	69		1		10
5(3)	1.5		14.5	5	62.5	3	0.5	10	
6(1)		4	18	7	50	5		15	
7(3)	1	1	12	4	64.5	2	0.5		12
8(2.5)		2.5	15	7.5	56.5		1	15	
9(1)			10	5	74.2		0.8		7
10(2)	2		10	3.5	67.5	5			10
11(2)		4	15	5	63.3		0.7	10	

TABLE V

Ex	AB9	RB15	PAA	X	W	BE	GBL	SURF	SDBS	AM
12(5)	2		13	4	55.2	5			0.8	15
13(3)		2	15	5	59		5		1	10
14(1)		2	8	4	74.5	2.5	2.5	0.5		5
15(0.5)	2		10	3	69.5	4		0.5	0.5	10
16(2)		2	15	7.5	52.5				1	20

TABLE VI

Ex	DB87	RB15	PAA	X	W	HT	HMSB	ADBS	AM
17(0.25)	3.5		13.5	4	63.5	1.75		0.5	14
18(0.1)		2.9	15	5	60	1		1	15
E (0.5)	3.5	0.5	12	6	65.5				12
1(0.25)	4.75	0.25	12	5	66.9			0.85	10
2(2)		0.5	12.5	4	68		1		12
3(1)		1.5	15	4.5	65.5	1	0.5	1	10
4(0.3)	3		12	4	67.8			0.9	12

TABLE VII

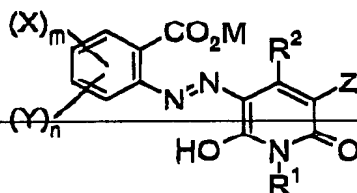
Ex	CD	PAA	X	TEA	W	HT	HMSB	NMP	BE	ADBS	AM
1(1)	3	15	5		53.5	2		5		0.5	15
2(2)	3	12		5	64	1	1		2		10
3(0.5)	4.5	12	2	2	64		2.5	2	2	0.5	8
4(0.2)	2.3	13	4.5		63.5	1.5				1	14

SMC 60384

31

CLAIMS

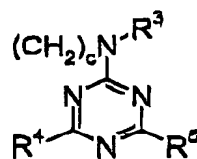
1. A compound of Formula 1



Formula 1

in which:

R^1 represents H, an optionally substituted C_{1-8} carbonyl derived group, or a group of Formula A



Formula A

where:

c is from 2 to 6,

R^3 represent H or optionally substituted C_{1-8} carbonyl derived group; and

R^4 and R^6 independently represent an optional substituent;

R^2 represents an optionally substituted C_{1-8} carbonyl derived group,

X , Y and Z independently represent H, an optional substituent or optionally substituted C_{1-8} carbonyl derived group;

M represents H or a cation

m and n independently represent 0, 1 or 2;

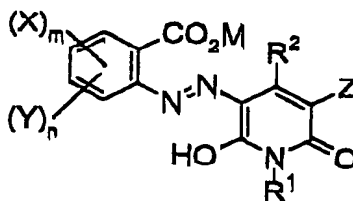
with the provisos that:

at least one of R^1 , R^2 , X , Y or Z comprises a group of formula SO_3M or PO_3M_2 where M is independently as represented herein;

when n is 0; m is 1; X is a sulfo group para to the azo group; Z is H and R^2 is methyl then R^1 is other than ethyl;

and the compound of formula 1 is other than a compound of formula II, III or IV as described herein.

2. A composition which comprises an ink effective for ink-jet printing and/or effective for use in a colour filter, the composition comprising a solvent and at least one compound of Formula 2;

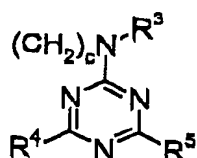


Formula 2

in which:

R^1 represents H, an optionally substituted C_{1-8} carbonyl derived group, or a group of

Formula A



Formula A

where:

c is from 2 to 6,

R^3 represent H or optionally substituted C_{1-8} carbonyl derived group; and

R^4 and R^5 independently represent an optional substituent;

R^2 represents an optionally substituted C_{1-8} carbonyl derived group,

X, Y and Z independently represent H, an optional substituent or optionally substituted C_{1-8} carbonyl derived group;

M represents H or a cation

m and n independently represent 0, 1 or 2.

3. A composition comprising a solvent and one or more dyes of Formula 1 as claimed in claim 1.

4. A composition as claimed in claim 2 or 3, wherein the solvent comprises water and one or more water soluble organic solvent(s)

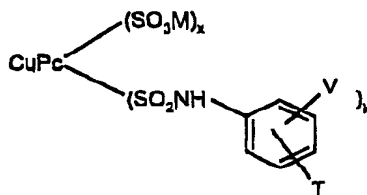
5. A composition as claimed in any of claims 2 to 4 which comprises at least one further colorant.

6. A composition as claimed in claim 5, where the further colorant(s) is selected from at least one cyan, green, red, magenta and/or orange colorant which is a dye or a pigment.

SMC 60384

33

7. A composition as claimed in claim 6, where the cyan dye is a compound of Formula (3)



Formula 3

in which:

T independently represents the same as given for X or Y in compounds of formula

1,

V represents CO_2M , SO_3M or PO_3M_2

x and y independently represent from 0 to 4; and

x + y is from 3 to 5,

where M is as given herein for compounds of formula 1.

8. A composition according to claim 6, where the cyan pigment is selected from C.I. Pigment Green 7 and C.I. Pigment Green 36.

9. A composition according to any of claims 2 to 8 which is a green, red or yellow ink suitable for use in any of the process claimed in claims 11 to 16.

10. Use of at least one compound of Formula 1 as claimed in claim 1, and/or a composition as claimed in any of claims 2 to 9, as a colorant for an ink jet printing ink.

11. A process for preparing a patterned, cross-linked, polymer film coating on a substrate; the process comprising the steps of:

(a) applying to the substrate simultaneously and/or sequentially in any order:

(i) one or more cross-linkable precursor(s) for one or more cross-linked polymer(s);

(ii) optionally one or more separate cross-linker(s) capable of cross-linking the precursor(s) for the polymer(s); and

(iii) one or more colorant(s) of Formula (1) as claimed in claim 1 or as represented in any of claims 2 to 8;

(b) optionally patterning one or more non cross-linked film(s) of component (i); component (ii); component (iii) and/or mixture(s) thereof, optionally before application of further components; and

(c) initiating cross-linking the mixture of components (i); (ii) in situ, to form an optionally patterned, cross-linked polymeric film coating on the substrate.

12. A process according to claim 11, in which the process is a printing process.

13. A process according to claim 12, in which the printing process is an ink-jet printing process.

14. A process according to claim 11, in which the process is a photolithographic process.

15. A process according to claim 11, in which the polymer precursor(s) comprise water dissolvable polymer precursor(s).

16. A process according to claim 15, in which the water dissolvable polymer precursor(s) comprise acrylic polymer precursor(s).

17. A substrate obtainable by a process as claimed in any of claims 11 to 16.

18. A substrate as claimed in claim 17, which comprises: a colour filter comprising a coloured, cross-linked, polymer coating on a transparent substrate; and/or a transparent, coloured, cross-linked, polymer coating on a substrate.

19. A substrate as claimed in either claim 17 or 18, which has utility as a component for a coloured display.

20. A substrate as claimed in any of claims 17 to 19, which comprises an array of coloured trichromatic elements in which the trichromat is selected from: a red, green and blue trichromat; and a cyan, magenta and yellow trichromat.

21. A display which comprises a substrate as claimed in any of claims 17 to 20.

22. A display as claimed in claim 21, which comprises a liquid crystal display.

23. An ink jet printer, component of said printer and/or consumable for use with said printer, which comprises a compound of Formula 1 as represented in claim 1; composition as claimed in any of claims 2 to 9, a substrate as claimed in any of claims 17 to 20, and/or a display as claimed in either claim 21 or 22.

24. An ink-jet printer, component and/or consumable as claimed in claim 23, comprising an ink-jet printer cartridge and/or replaceable ink-jet printing head.

SMC 60384

35

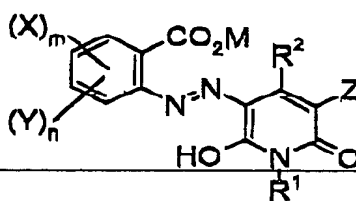
25. A process, composition, ink, substrate, display, Ink-jet printer, component of said printer and/or consumable for use with said printer, which is substantially as described herein, with reference to the examples.

5 26. A process for printing a substrate with a composition according to any of claims 2 to 9 using an ink-jet printer.

27. A paper, overhead projector slide, textile or colour filter printed with a process according to claim 26.

ABSTRACT

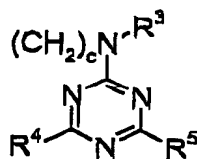
There is described compounds of Formula 1:



Formula 1

In which:

R^1 represents H, an optionally substituted C_{1-8} carbonyl derived group, or a group of Formula A



Formula A

where:

c is from 2 to 6,

R^3 represent optionally substituted C_{1-8} carbonyl derived group; and

R^4 and R^5 independently represent an optionally substituent;

R^2 represents an optionally substituted C_{1-8} carbonyl derived group,

X , Y and Z independently represent H, an optional substituent or optionally substituted C_{1-8} carbonyl derived group;

M represents H or a cation

m and n independently represent 0, 1 or 2;

at least one of R^1 , R^2 , X , Y or Z comprises a group of formula SO_3M or PO_3M_2 where M is independently as represented herein.

These compounds are useful as the colorants to prepare colour filters for displays.